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January 30, 2002

FILE: UTSC:671US

CERTIFICATE OF MAILING 37 C.F.R 1.8

I hereby certify that this correspondence is being deposited with the U.S. Postal Service with sufficient postage as First Class Mail in an envelope addressed to: Commissioner for Patents, Washington, DC 20231, on the date below:

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Gina N. Shishima

Commissioner for Patents Washington, DC 20231

RE:

SN 09/978,318 "WWOX: A TUMOR SUPPRESSOR GENE MUTATED IN MULTIPLE

CANCERS" - C. Marcelo Aldaz and Andrzej Bednarek

Sir:

Enclosed for filing in the above-referenced patent application is an Information Disclosure Statement, Form PTO-1449, and references (B1-B2, C1-C46).

No fees are believed to be due in connection with the filing of this Information Disclosure Statement, however, should any fees under 37 C.F.R. §§ 1.16 to 1.21 be deemed necessary for any reason relating to the enclosed materials, the Commissioner is hereby authorized to deduct said fees from Fulbright & Jaworski Deposit Account No.: 50-1212/10109289/GNS.

Please date stamp and return the enclosed postcard evidencing receipt of these materials.

Respectfully submitted,

Øina N. Shishima Reg. No. 45,104

GNS/cmb

Encl: as noted

25114553.1





N THE UNITED STATES PATENT AND TRADEMARK OFFICE

C. Marcelo Aldaz Andrzej Bednarek

Serial No.: 09/978,318

Filed: October 15, 2001

For: WWOX: A TUMOR SUPPRESSOR GENE

MUTATED IN MULTIPLE CANCERS

Group Art Unit: Unknown

Examiner: Unknown

Atty. Dkt. No.: UTSC:671US/GNS

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Gina N. Shishima

INFORMATION DISCLOSURE STATEMENT

Commissioner for Patents Washington, D.C. 20231

Sir:

In compliance with the duty of disclosure under 37 C.F.R. § 1.56, it is respectfully requested that this Information Disclosure Statement be entered and the documents listed on attached Form PTO-1449 be considered by the Examiner and made of record. Copies of the listed documents required by 37 C.F.R. § 1.98(a)(2) are enclosed for the convenience of the Examiner.

In accordance with 37 C.F.R §§ 1.97(g), (h), this Information Disclosure Statement is not to be construed as a representation that a search has been made, and is not to be construed to be an admission that the information cited is, or is considered to be, material to patentability as defined in 37 C.F.R. § 1.56(b).

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The present Information Disclosure Statement is being filed prior to the receipt of a first Official Action reflecting an examination on the merits, and hence is believed to be timely filed in accordance with 37 C.F.R § 1.97(b). No fees are believed to be due in connection with the filing of this Information Disclosure Statement, however, should any fees under 37 C.F.R. §§ 1.16 to 1.21 be deemed necessary for any reason relating to these materials, the Commissioner is hereby authorized to deduct said fees from Fulbright & Jaworski Deposit Account No.: 50-1212/10109289/GNS.

Applicants respectfully request that the listed documents be made of record in the present case.

Respectfully submitted,

Gina N. Shishima

Reg. No. 45,104

Attorney for Applicants

FULBRIGHT & JAWORSKI L.L.P. 600 Congress Avenue, Suite 2400 Austin, Texas 78701 (512) 474-5201

Date:

January 30, 2002

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	C1	Aldaz et al., "Comparative allelotype of in situ and invasive human breast cancer: high frequency of microsatellite instability in lobular breast carcinomas," Cancer Res., 55:3976-3981, 1995.								
	C2	Andre and Springael, "WWP, a new amino acid motif present in single or multiple copies in various proteins including dystrophin and the SH3-binding Yes- associated protein YAP65," <i>Biochem. Biophys. Res. Commun.</i> , 205(2):1201-1205, 1994.								
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	C4	Bednarek and Al chromosome 16 39:128, #872, 19	(q23.3-q24.1):	rizatio in hum	n of transcrip an breast can	ts fro	om a coi Proc. A	mmonly de Imer. Asso	eleted area of oc. Cancer Res.,	
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List of Patents and Publications for INFORMATION DISCLOSUREM	Applicate s MEMENT	Applicant C. Marcelo Aldaz Andrzej Bednarek		
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U.S. Patent Documents	Foreign	Patent Documents	Other Art	

U.S. Patent Documents	Foreign Patent Documents	Other Art
See Page 1	See Page 1	See Page 1

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	C7	Bork and Sudol, "The WW domain: a signalling site in dystrophin?" <i>Trends Biochem. Sci.</i> , 19:531-533, 1994.
	C8	Carter et al., "Allelic loss of chromosomes 16q and 10q in human prostate cancer," Proc. Natl. Acad. Sci. USA, 87: 8751-8755, 1990
	C9	Chan et al., "Formin binding proteins bear WWP/WW domains that bind proline-rich peptides and functionally resembel SH3 domains," EMBO J., 15(5):1045-1054, 1996.
	C10	Chang et al., "Hyaluronidase induction of a WW domain-containing oxidoreductase that enhanced tumor necrosis factor cytotoxicity," J. Biol. Chem., 276:3361-3370, 2001.
	C11	Chen and Sudol, "The WW domain of Yes-associated protein binds a proline-rich ligand that differs from the consensus established for Src homology 3-binding modules," <i>Proc. Natl. Acad. Sci. USA</i> , 92:7819-7823, 1995.
	C12	Chen et al., "Deletion map of chromosome 16q in ductal carcinoma in situ of the breast: refining a putative tumor suppressor gene region," Cancer Res. 56:5605-5609, 1996.
	C13	Chesi et al., "Frequent dysregulation of the c-maf proto-oncogene at 16q23 by translocation to an Ig locus in multiple myeloma," Blood, 91:4457-4463, 1998.
	C14	Cleton-Jansen et al., "At least two different regions are involved in allelic imbalance on chromosome arm 16q in breast cancer," Genes, Chromos, Cancer, 9:101-107, 1994.
	C15	Crawford et al., "The PISSLRE gene: structure, exon skipping, and exclusion as tumor suppressor in breast cancer," Genomics, 56:90-97, 1999.
	C16	Duax and Ghosh, "Structure and function of steroid dehydrogenases involved in hypertension, fertility, and cancer," <i>Steroids</i> , 62:95-100, 1997.
	C17	Dutrillaux et al., "Characterization of chromosomal anomalies in human breast cancer. A comparison of 30 paradiploid cases with few chromosome changes," Cancer Genet. Cytogenet., 49:203-217, 1990.
	C18	GenBank Accession Number AF179633
	C19	GenBank Accession Number AF211943
	C20	GenBank Accession Number AF212843
	C21	GenBank Accession Number AF227526

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EXAMINER: DATE CONSIDERED:

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1	7 2002 & UTSO	Docket No. C:671US/GNS	Serial No. 09/978,318	
List of Patents and Publications for Information Disclosure S	Applicant's Applicant C. M. C. M. Andrews Andrews Andrews Andrews Andrews Andrews Applicant Appl	icant arcelo Aldaz zej Bednarek		
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	C22	GenBank Accession Number AF227527
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	C24	GenBank Accession Number AF395123
	C25	GenBank Accession Number AF395124
	C26	GenBank Accession Number U13395, locus ID 9621
	C27	Jornvall et al., "Short-chain dehydrogenases/reductases (SDR)," Biochemistry, 34:6003-6013, 1995.
	C28	Krummel et al., "The characterization of the common fragile site FRA16D and its involvement in multiple myeloma translocations," Genomics, 69:37-46, 2000.
	C29	Lu et al., "Function of WW domains as phosphoserine- or phosphothreonine-binding modules," Science, 283:1325-1328, 1999.
	C30	Mangelsdorf et al., "Chromosomal fragile site FRA16D and DNA instability in cancer," Cancer Res., 60: 1683-1689, 2000.
	C31	Paige et al., "A 700-kb physical map of a region of 16q23.2 homozygously deleted in multiple cancers and spanning the common fragile site FRA16D," Cancer Res. 60:1690-1697, 2000.
	C32	Paige et al., "WWOX: A candidate tumor suppressor gene involved in multiple tumor types," Proc. Natl. Acad. Sci. USA, 98:11417-11422, 2001
	C33	Pandis et al., "Whole-arm t(1;16) and i(1q) as sole anomalies identify gain of 1q as a primary chromosomal abnormality in breast cancer," Genes Chromosomes Cancer, 5:235-238, 1992.
	C34	Price et al., "Tumorigenicity and metastasis of human breast carcinoma cell lines in nude mice," Cancer Res. 50:717-721, 1990.
	C35	Richards, "Fragile and unstable chromosomes in cancer: causes and consequences," <i>Trends Genet.</i> , 17:339-345, 2001.
	C36	Ried et al., "Common chormosomal fragile site FRA16D sequence: identification of the FOR gene spanning FRA16D and homozygous deletions and translocation breakpoints in cancer cells,""Human Molecular Genetics, 9(11):1651-1663, 2000.

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Atty. Docket No. Serial No. Form PTO-1449 (modified) FEB 0 7 2002 UTSC:671US/GNS 09/978,318 Applicant List of Patents and Publications for Applican C. Marcelo Aldaz Andrzej Bednarek INFORMATION DISCLOSUR Filing Date: Group: (Use several sheets if necessary) October 15, 2001 Unknown **Foreign Patent Documents** Other Art **U.S. Patent Documents** See Page 1 See Page 1 See Page 1

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	C37	Sato et al., "Allelotype of breast cancer: cumulative allele losses promote tumor progression in primary breast cancer," Cancer Res., 50:7184-7189, 1990.
	C38	Savino et al., "Characterization of copine VII, a new member of the copine family, and its exclusion as a candidate in sporadic breast cancers with loss of heterozygosity at 16q24.3," Genomics, 61:219-226, 1999.
	C39	Smith et al., "Common fragile sites and cancer (Review)," Int. J. Oncol., 12:187-196, 1998.
	C40	Staub et al., "WW domains of Nedd4 bind to the proline-rich PY motifs in the epithelial Na+channel deleted in Liddle's syndrome," Embo. J., 15:2371-2380, 1996.
	C41	Sudol and Hunter, "NeW wrinkles for an old domain," Cell, 103:1001-1004, 2000.
	C42	Sudol et al., "Characterization of the mammalian YAP (Yes-associated protein) gene and its role in defining a novel protein module, the WW domain," J. Biol. Chem., 270:14733-14741, 1995.
,	C43	Sudol, "Yes-associated protein (YAP65) is a proline-rich phosphoprotein that binds to the SH3 domain of the Yes proto-oncogene product," <i>Oncogene</i> , 9:2145-2152, 1994.
	C44	Sutherland et al., "Fragile sites still breaking," Trends Genet., 14:501-506, 1998.
	C45	Tsuda et al., "Allele loss on chromosome 16q242-qter occurs frequently in breast cancer irrespectively of differences in phenotype and extent of spread," Cancer Res., 54: 513-517, 1994.
	C46	Whitmore et al., "Construction of a high-resolution physical and transcription map of chromosome 16q24.3: a region of frequent loss of heterozygosity in sporadic breast cancer," Genomics, 50:1-8, 1998.

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